

To: Office of Attorney General
AGO.highcostprescriptiondrugs@vermont.gov

From: Mylan Pharmaceuticals Inc.
 781 Chestnut Ridge Road
 Morgantown, WV 26505

Date: October 9, 2020

Re: 18 V.S.A § 4637

In compliance with 18 V.S.A. § 4637, on September 11, 2020 Mylan Pharmaceuticals Inc. ("Mylan") provided written notice to the Office of the Attorney General that it introduced a new prescription drug, Everolimus Tablets ("the Product"), to the commercial market on September 10, 2020 at a wholesale acquisition cost that is over the threshold set for a specialty drug under the Medicare Part D program.

This letter provides the additional required information by 18 V.S.A. § 4637(c) regarding the Product. Mylan notes that the Office of the Attorney General has not yet prescribed a format for submissions under this section. Further, as authorized by 18 V.S.A. § 4637(d), Mylan has limited the information reported to that which is in the public domain or publicly available.

- (1) A description of the marketing and pricing plans used in the launch of the new drug in the United States and internationally;

Everolimus tablets are indicated (1) for the treatment postmenopausal women with advanced hormone receptor-positive, HER2-negative breast cancer in combination with exemestane, after failure of treatment with letrozole or anastrozole; (2) for the treatment of adult patients with renal angiomyolipoma and Tuberous Sclerosis Complex (TSC), not requiring immediate surgery; and (3) in adult and pediatric patients aged 1 year and older with Tuberous Sclerosis Complex (TSC) for the treatment of Subependymal Giant Cell Astrocytoma (SEGA) that requires therapeutic intervention but cannot be curatively resected. The Wholesale Acquisition Cost (WAC) for the product in the United States is below:

NDC	Product	Package Size	WAC
00378-3096-85	Everolimus Tablets, 2.5mg	28	\$6,419.43
00378-3097-85	Everolimus Tablets, 5mg	28	\$6,714.62
00378-3098-85	Everolimus Tablets, 7.5mg	28	\$6,714.62

The prices negotiated with customers as well as any marketing plans in the United States or internationally are confidential and not in the public domain or publicly available. In the United States, Mylan sells its products directly to wholesalers, distributors, retail pharmacy chains, long-term care facilities and mail order pharmacies. Mylan also sells its generic products indirectly to several entities, including independent pharmacies, managed care organizations, hospitals, etc. These customers, called "indirect customers," purchase our products primarily through our wholesale customers.

(2) the estimated volume of patients who may be prescribed the drug;

No information specific to the estimated number of patients that may be prescribed Mylan's Product is in the public domain or publicly available.

(3) whether the drug was granted breakthrough therapy designation or priority review by the FDA prior to final approval;

The Product was not granted breakthrough therapy designation or priority review by the FDA.

(4) the date and price of acquisition if the drug was not developed by the manufacturer.

The Product was not the result of an acquisition.